

**BEFORE THE
MEDICAL BOARD OF CALIFORNIA
DEPARTMENT OF CONSUMER AFFAIRS
STATE OF CALIFORNIA**

**In the Matter of the Accusation
Against:**

Moshen T. Moghaddam, M.D.

**Physician's and Surgeon's
Certificate No. A 46373**

Respondent.

Case No. 800-2018-042453

DECISION

**The attached Stipulated Surrender of License and Order is hereby
adopted as the Decision and Order of the Medical Board of California,
Department of Consumer Affairs, State of California.**

This Decision shall become effective at 5:00 p.m. on APR 21 2022.

IT IS SO ORDERED APR 14 2022.

MEDICAL BOARD OF CALIFORNIA


for: **William Prasifka** **Reji Varghese**
Executive Director **Deputy Director**

1 ROB BONTA
Attorney General of California
2 ROBERT MCKIM BELL
Supervising Deputy Attorney General
3 VLADIMIR SHALKEVICH
Deputy Attorney General
4 State Bar No. 173955
300 So. Spring Street, Suite 1702
5 Los Angeles, California 90013
Telephone: (213) 269-6538
6 Facsimile: (916) 731-2117
Attorneys for Complainant
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8 **BEFORE THE**
9 **MEDICAL BOARD OF CALIFORNIA**
10 **DEPARTMENT OF CONSUMER AFFAIRS**
11 **STATE OF CALIFORNIA**

12 In the Matter of the Accusation Against:

Case No. 800-2018-042453

13 MOSHEN T. MOGHADDAM, M.D.

14 19100 Ventura Boulevard, Suite 16
Tarzana, California 91356-3234

15 Physician's and Surgeon's Certificate A 46373,
16 Respondent.

**STIPULATED SURRENDER OF
LICENSE AND ORDER**

17
18 **IT IS HEREBY STIPULATED AND AGREED** by and between the parties to the above-
19 entitled proceedings that the following matters are true:

20 **PARTIES**

21 1. William Prasifka (Complainant) is the Executive Director of the Medical Board of
22 California (Board). He brought this action solely in his official capacity and is represented in this
23 matter by Rob Bonta, Attorney General of the State of California, by Vladimir Shalkevich,
24 Deputy Attorney General.

25 2. Moshen T. Moghaddam, M.D. (Respondent) is represented in this proceeding by
26 attorney Peter R. Osinoff of Bonne, Bridges, Mueller, O'Keefe & Nichols, 355 South Grand
27 Avenue, Suite 1750, Los Angeles, California 90071-1562.

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3. On August 7, 1989, the Board issued Physician's and Surgeon's Certificate No. A 46373 to Moshen T. Moghaddam, M.D. (Respondent). That license was in full force and effect at all times relevant to the charges brought in Accusation No. 800-2018-042453 and will expire on December 31, 2022, unless renewed.

JURISDICTION

4. Accusation No. 800-2018-042453 was filed before the Board and is currently pending against Respondent. The Accusation and all other statutorily required documents were properly served on Respondent on March 24, 2021. Respondent timely filed his Notice of Defense contesting the Accusation. A copy of Accusation No. 800-2018-042453 is attached as Exhibit A and is incorporated by reference.

ADVISEMENT AND WAIVERS

5. In addition to the Accusation No. 800-2018-042453, the Board is currently conducting two additional investigations of complaints pertaining to Respondent, case numbers 800-2019-054896 and 800-2021-078617. Respondent has no prior record of discipline and wishes to retire from the practice of medicine. It is the intent of the parties that this Stipulated Surrender will resolve all disciplinary charges pending against Respondent as well as any potential disciplinary charges represented by the ongoing investigations.

6. Respondent has carefully read, fully discussed with counsel, and understands the charges and allegations in Accusation No. 800-2018-042453. Respondent also has carefully read, fully discussed with counsel, and understands the effects of this Stipulated Surrender of License and Order.

7. Respondent is fully aware of his legal rights in this matter, including the right to a hearing on the charges and allegations in the Accusation; the right to confront and cross-examine the witnesses against him; the right to present evidence and to testify on his own behalf; the right to the issuance of subpoenas to compel the attendance of witnesses and the production of documents; the right to reconsideration and court review of an adverse decision; and all other rights accorded by the California Administrative Procedure Act and other applicable laws.

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8. Respondent voluntarily, knowingly, and intelligently waives and gives up each and every right set forth above.

CULPABILITY

9. Respondent understands that the charges and allegations in Accusation No. 800-2018-042453, if proven at a hearing, constitute cause for imposing discipline upon his Physician's and Surgeon's Certificate.

10. For the purpose of resolving the Accusation without the expense and uncertainty of further proceedings, Respondent agrees that, at a hearing, Complainant could establish a factual basis for the charges in the Accusation and that those charges constitute cause for discipline. Respondent hereby gives up his right to contest that cause for discipline exists based on those charges.

11. Respondent understands that by signing this stipulation he enables the Board to issue an order accepting the surrender of his Physician's and Surgeon's Certificate without further process.

CONTINGENCY

12. This stipulation shall be subject to approval by the Board. Respondent understands and agrees that counsel for Complainant and the staff of the Board may communicate directly with the Board regarding this stipulation and surrender, without notice to or participation by Respondent or his counsel. By signing the stipulation, Respondent understands and agrees that he may not withdraw his agreement or seek to rescind the stipulation prior to the time the Board considers and acts upon it. If the Board fails to adopt this stipulation as its Decision and Order, the Stipulated Surrender and Disciplinary Order shall be of no force or effect, except for this paragraph, it shall be inadmissible in any legal action between the parties, and the Board shall not be disqualified from further action by having considered this matter.

13. The parties understand and agree that Portable Document Format (PDF) and facsimile copies of this Stipulated Surrender of License and Order, including PDF and facsimile signatures thereto, shall have the same force and effect as the originals.

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14. In consideration of the foregoing admissions and stipulations, the parties agree that the Board may, without further notice or formal proceeding, issue and enter the following Order:

ORDER

IT IS HEREBY ORDERED THAT Physician's and Surgeon's Certificate No. A 46373,
issued to Respondent Moshen T. Moghaddam, M.D., is surrendered and accepted by the Board.

1. The surrender of Respondent's Physician's and Surgeon's Certificate and the acceptance of the surrendered license by the Board shall constitute the imposition of discipline against Respondent. This stipulation constitutes a record of the discipline and shall become a part of Respondent's license history with the Board.

2. Respondent shall lose all rights and privileges as a Physician and Surgeon in California as of the effective date of the Board's Decision and Order.

3. Respondent shall cause to be delivered to the Board his pocket license and, if one was issued, his wall certificate on or before the effective date of the Decision and Order.

4. If Respondent ever files an application for licensure or a petition for reinstatement in the State of California, the Board shall treat it as a petition for reinstatement. Respondent must comply with all the laws, regulations and procedures for reinstatement of a revoked or surrendered license in effect at the time the petition is filed, and all of the charges and allegations contained in Accusation No. 800-2018-042453 shall be deemed to be true, correct and admitted by Respondent when the Board determines whether to grant or deny the petition.

5. In addition, and notwithstanding any applicable period of limitations, if and/or when the Board determines whether to grant or deny any future petition for reinstatement by Respondent, the Board shall consider the circumstances of ongoing investigations 800-2019-054896 and 800-2021-078617, and admit into evidence in any reinstatement proceeding reports of investigation number 800-2019-054896 and 800-2021-078617 and any attachments thereto.

6. Respondent shall pay the agency its costs of investigation and enforcement in the amount of \$3,625 prior to issuance of a new or reinstated license.

7. If Respondent should ever apply or reapply for a new license or certification, or petition for reinstatement of a license, by any other health care licensing agency in the State of

1 California, all of the charges and allegations contained in Accusation, No. 800-2018-042453 shall
2 be deemed to be true, correct, and admitted by Respondent for the purpose of any Statement of
3 Issues or any other proceeding seeking to deny or restrict licensure. The circumstances of the
4 ongoing investigations 800-2019-054896 and 800-2021-078617, shall also be considered, and
5 reports of investigation number 800-2019-054896 and 800-2021-078617, including any
6 attachments thereto, shall be admitted into evidence of any Statement of Issues or any other
7 proceeding seeking to deny or restrict licensure.

8 **ACCEPTANCE**

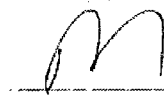
9 I have carefully read the above Stipulated Surrender of License and Order and have fully
10 discussed it with my attorney Peter R. Osinoff, Esq. I understand the stipulation and the effect it
11 will have on my Physician's and Surgeon's Certificate. I enter into this Stipulated Surrender of
12 License and Order voluntarily, knowingly, and intelligently, and agree to be bound by the
13 Decision and Order of the Medical Board of California.

14
15 DATED: 4.3.2022


MOSHEN T. MOGHADDAM, M.D.
Respondent

17 I have read and fully discussed with Respondent MOSHEN T. MOGHADDAM, M.D. the
18 terms and conditions and other matters contained in this Stipulated Surrender of License and
19 Order. I approve its form and content.

20 DATED: 4/6/2022


PETER R. OSINOFF
Attorney for Respondent

1 **ENDORSEMENT**

2 The foregoing Stipulated Surrender of License and Order is hereby respectfully submitted
3 for consideration by the Medical Board of California of the Department of Consumer Affairs.

4 DATED: April 7, 2022

Respectfully submitted,

5 ROB BONTA
6 Attorney General of California
7 ROBERT MCKIM BELL
8 Supervising Deputy Attorney General

9  Electronically Signed

10 VLADIMIR SHALKEVICH
11 Deputy Attorney General
12 *Attorneys for Complainant*

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14 64988590.docx

Exhibit A

Accusation No. 800-2018-042453

1 MATTHEW RODRIQUEZ
Acting Attorney General of California
2 ROBERT MCKIM BELL
Supervising Deputy Attorney General
3 COLLEEN M. MCGURRIN
Deputy Attorney General
4 California Department of Justice
State Bar No. 147250
5 300 South Spring Street, Suite 1702
Los Angeles, California 90013
6 Telephone: (213) 269-6546
Facsimile: (916) 731-2117
7 *Attorneys for Complainant*

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Case No. 800-2018-042453

13 MOSHEN T. MOGHADDAM, M.D.

A C C U S A T I O N

14 19100 Ventura Boulevard, Suite 16
Tarzana, California 91356-3234

15 Physician's and Surgeon's Certificate A 46373,

16 Respondent.

17 **PARTIES**

18
19 1. William Prasifka (Complainant) brings this Accusation solely in his official capacity
20 as the Executive Director of the Medical Board of California, Department of Consumer Affairs
21 (Board).

22 2. On or about August 7, 1989, the Board issued Physician's and Surgeon's Certificate
23 Number A 46373 to Moshen T. Moghaddam, M.D. (Respondent). That license was in full force
24 and effect at all times relevant to the charges brought herein and will expire on December 31,
25 2022, unless renewed.

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4. Section 2227 of the Code states:

(1) Have his or her license revoked upon order of the board.

(4) Be publicly reprimanded by the board. The public reprimand may include a requirement that the licensee complete relevant educational courses approved by the board.

(b) Any matter heard pursuant to subdivision (a), except for warning letters, medical review or advisory conferences, professional competency examinations, continuing education activities, and cost reimbursement associated therewith that are agreed to with the board and successfully completed by the licensee, or other matters made confidential or privileged by existing law, is deemed public, and shall be made available to the public by the board pursuant to Section 803.1.

5. Section 2234 of the Code states:

(b) Gross negligence.

(c) **Repeated negligent acts.** To be repeated, there must be two or more negligent acts or omissions. An initial negligent act or omission followed by a separate and distinct departure from the applicable standard of care shall constitute repeated negligent acts.

1 (1) An initial negligent diagnosis followed by an act or omission medically
2 appropriate for that negligent diagnosis of the patient shall constitute a single
3 negligent act.

4 (2) When the standard of care requires a change in the diagnosis, act, or
5 omission that constitutes the negligent act described in paragraph (1), including, but
6 not limited to, a reevaluation of the diagnosis or a change in treatment, and the
7 licensee's conduct departs from the applicable standard of care, each departure
8 constitutes a separate and distinct breach of the standard of care.

9 (d) Incompetence.

10 (e) The commission of any act involving dishonesty or corruption that is
11 substantially related to the qualifications, functions, or duties of a physician and
12 surgeon.

13 (f) Any action or conduct that would have warranted the denial of a certificate.

14 (g) The failure by a certificate holder, in the absence of good cause, to attend
15 and participate in an interview by the board. This subdivision shall only apply to a
16 certificate holder who is the subject of an investigation by the board.

17 6. Section 2220 of the Code states:

18 Except as otherwise provided by law, the board may take action against all
19 persons guilty of violating this chapter. The board shall enforce and administer this
20 article as to physician and surgeon certificate holders, including those who hold
21 certificates that do not permit them to practice medicine, such as, but not limited to,
22 retired, inactive, or disabled status certificate holders, and the board shall have all the
23 powers granted in this chapter for these purposes including, but not limited to:

24 (a) Investigating complaints from the public, from other licensees, from health
25 care facilities, or from the board that a physician and surgeon may be guilty of
26 unprofessional conduct. The board shall investigate the circumstances underlying a
27 report received pursuant to Section 805 or 805.01 within 30 days to determine if an
28 interim suspension order or temporary restraining order should be issued. The board
shall otherwise provide timely disposition of the reports received pursuant to Section
805 and Section 805.01.

(b) Investigating the circumstances of practice of any physician and surgeon
where there have been any judgments, settlements, or arbitration awards requiring the
physician and surgeon or his or her professional liability insurer to pay an amount in
damages in excess of a cumulative total of thirty thousand dollars (\$30,000) with
respect to any claim that injury or damage was proximately caused by the physician's
and surgeon's error, negligence, or omission.

(c) Investigating the nature and causes of injuries from cases which shall be
reported of a high number of judgments, settlements, or arbitration awards against a
physician and surgeon.

7. Section 2228 of the Code states:

The authority of the board or the California Board of Podiatric Medicine to
discipline a licensee by placing him or her on probation includes, but is not limited to,
the following:

1 (a) Requiring the licensee to obtain additional professional training and to pass
2 an examination upon the completion of the training. The examination may be written
or oral, or both, and may be a practical or clinical examination, or both, at the option
of the board or the administrative law judge.

3 (b) Requiring the licensee to submit to a complete diagnostic examination by
4 one or more physicians and surgeons appointed by the board. If an examination is
5 ordered, the board shall receive and consider any other report of a complete
diagnostic examination given by one or more physicians and surgeons of the
licensee's choice.

6 (c) Restricting or limiting the extent, scope, or type of practice of the licensee,
7 including requiring notice to applicable patients that the licensee is unable to perform
the indicated treatment, where appropriate.

8 (d) Providing the option of alternative community service in cases other than
9 violations relating to quality of care.

10 8. Section 2241.5 of the Code states:

11 (a) A physician and surgeon may prescribe for, or dispense or administer to, a
12 person under his or her treatment for a medical condition dangerous drugs or
prescription controlled substances for the treatment of pain or a condition causing
pain, including, but not limited to, intractable pain.

13 (b) No physician and surgeon shall be subject to disciplinary action for
14 prescribing, dispensing, or administering dangerous drugs or prescription controlled
substances in accordance with this section.

15 (c) This section shall not affect the power of the board to take any action
16 described in Section 2227 against a physician and surgeon who does any of the
following:

17 (1) Violates subdivision (b), (c), or (d) of Section 2234 regarding gross
18 negligence, repeated negligent acts, or incompetence.

19 (2) Violates Section 2241 regarding treatment of an addict.

20 (3) Violates Section 2242 or 2525.3 regarding performing an appropriate prior
21 examination and the existence of a medical indication for prescribing, dispensing, or
furnishing dangerous drugs or recommending medical cannabis.

22 (4) Violates Section 2242.1 regarding prescribing on the Internet.

23 (5) Fails to keep complete and accurate records of purchases and disposals of
24 substances listed in the California Uniform Controlled Substances Act (Division 10
(commencing with Section 11000) of the Health and Safety Code) or controlled
25 substances scheduled in the federal Comprehensive Drug Abuse Prevention and
Control Act of 1970 (21 U.S.C. Sec. 801 et seq.), or pursuant to the federal
26 Comprehensive Drug Abuse Prevention and Control Act of 1970. A physician and
surgeon shall keep records of his or her purchases and disposals of these controlled
27 substances or dangerous drugs, including the date of purchase, the date and records of
the sale or disposal of the drugs by the physician and surgeon, the name and address
28 of the person receiving the drugs, and the reason for the disposal or the dispensing of
the drugs to the person, and shall otherwise comply with all state recordkeeping
requirements for controlled substances.

1 (6) Writes false or fictitious prescriptions for controlled substances listed in the
2 California Uniform Controlled Substances Act or scheduled in the federal
3 Comprehensive Drug Abuse Prevention and Control Act of 1970.

4 (7) Prescribes, administers, or dispenses in violation of this chapter, or in
5 violation of Chapter 4 (commencing with Section 11150) or Chapter 5 (commencing
6 with Section 11210) of Division 10 of the Health and Safety Code.

7 (d) A physician and surgeon shall exercise reasonable care in determining
8 whether a particular patient or condition, or the complexity of a patient's treatment,
9 including, but not limited to, a current or recent pattern of drug abuse, requires
10 consultation with, or referral to, a more qualified specialist.

11 (e) Nothing in this section shall prohibit the governing body of a hospital from
12 taking disciplinary actions against a physician and surgeon pursuant to Sections
13 809.05, 809.4, and 809.5.

14 9. Section 2242 of the Code states:

15 (a) Prescribing, dispensing, or furnishing dangerous drugs as defined in Section
16 4022 without an appropriate prior examination and a medical indication, constitutes
17 unprofessional conduct. An appropriate prior examination does not require a
18 synchronous interaction between the patient and the licensee and can be achieved
19 through the use of telehealth, including, but not limited to, a self-screening tool or a
20 questionnaire, provided that the licensee complies with the appropriate standard of
21 care.

22 (b) No licensee shall be found to have committed unprofessional conduct within
23 the meaning of this section if, at the time the drugs were prescribed, dispensed, or
24 furnished, any of the following applies:

25 (1) The licensee was a designated physician and surgeon or podiatrist serving in
26 the absence of the patient's physician and surgeon or podiatrist, as the case may be,
27 and if the drugs were prescribed, dispensed, or furnished only as necessary to
28 maintain the patient until the return of the patient's practitioner, but in any case no
longer than 72 hours.

(2) The licensee transmitted the order for the drugs to a registered nurse or to a
licensed vocational nurse in an inpatient facility, and if both of the following
conditions exist:

(A) The practitioner had consulted with the registered nurse or licensed
vocational nurse who had reviewed the patient's records.

(B) The practitioner was designated as the practitioner to serve in the absence
of the patient's physician and surgeon or podiatrist, as the case may be.

(3) The licensee was a designated practitioner serving in the absence of the
patient's physician and surgeon or podiatrist, as the case may be, and was in
possession of or had utilized the patient's records and ordered the renewal of a
medically indicated prescription for an amount not exceeding the original prescription
in strength or amount or for more than one refill.

(4) The licensee was acting in accordance with Section 120582 of the Health
and Safety Code.

10. Section 725 of the Code states:

(a) Repeated acts of clearly excessive prescribing, furnishing, dispensing, or administering of drugs or treatment, repeated acts of clearly excessive use of diagnostic procedures, or repeated acts of clearly excessive use of diagnostic or treatment facilities as determined by the standard of the community of licensees is unprofessional conduct for a physician and surgeon, dentist, podiatrist, psychologist, physical therapist, chiropractor, optometrist, speech-language pathologist, or audiologist.

(b) Any person who engages in repeated acts of clearly excessive prescribing or administering of drugs or treatment is guilty of a misdemeanor and shall be punished by a fine of not less than one hundred dollars (\$100) nor more than six hundred dollars (\$600), or by imprisonment for a term of not less than 60 days nor more than 180 days, or by both that fine and imprisonment.

(c) A practitioner who has a medical basis for prescribing, furnishing, dispensing, or administering dangerous drugs or prescription controlled substances shall not be subject to disciplinary action or prosecution under this section.

(d) No physician and surgeon shall be subject to disciplinary action pursuant to this section for treating intractable pain in compliance with Section 2241.5.

11. Section 2266 of the Code states: The failure of a physician and surgeon to maintain adequate and accurate records relating to the provision of services to their patients constitutes unprofessional conduct.

12. Health and Safety Code section 11165 states:

(a) To assist health care practitioners in their efforts to ensure appropriate prescribing, ordering, administering, furnishing, and dispensing of controlled substances, law enforcement and regulatory agencies in their efforts to control the diversion and resultant abuse of Schedule II, Schedule III, and Schedule IV controlled substances, and for statistical analysis, education, and research, the Department of Justice shall, contingent upon the availability of adequate funds in the CURES Fund, maintain the Controlled Substance Utilization Review and Evaluation System (CURES) for the electronic monitoring of, and Internet access to information regarding, the prescribing and dispensing of Schedule II, Schedule III, and Schedule IV controlled substances by all practitioners authorized to prescribe, order, administer, furnish, or dispense these controlled substances.

(b) The Department of Justice may seek and use grant funds to pay the costs incurred by the operation and maintenance of CURES. The department shall annually report to the Legislature and make available to the public the amount and source of funds it receives for support of CURES.

(c) (1) The operation of CURES shall comply with all applicable federal and state privacy and security laws and regulations.

(2) (A) CURES shall operate under existing provisions of law to safeguard the privacy and confidentiality of patients. Data obtained from CURES shall only be provided to appropriate state, local, and federal public agencies for disciplinary, civil, or criminal purposes and to other agencies or entities, as determined by the Department of Justice, for the purpose of educating practitioners and others in lieu of

1 disciplinary, civil, or criminal actions. Data may be provided to public or private
2 entities, as approved by the Department of Justice, for educational, peer review,
3 statistical, or research purposes, if patient information, including any information that
4 may identify the patient, is not compromised. Further, data disclosed to any
5 individual or agency as described in this subdivision shall not be disclosed, sold, or
6 transferred to any third party, unless authorized by, or pursuant to, state and federal
7 privacy and security laws and regulations. The Department of Justice shall establish
8 policies, procedures, and regulations regarding the use, access, evaluation,
9 management, implementation, operation, storage, disclosure, and security of the
10 information within CURES, consistent with this subdivision.

11 (B) Notwithstanding subparagraph (A), a regulatory board whose licensees do
12 not prescribe, order, administer, furnish, or dispense controlled substances shall not
13 be provided data obtained from CURES.

14 (3) The Department of Justice shall, no later than July 1, 2020, adopt
15 regulations regarding the access and use of the information within CURES. The
16 Department of Justice shall consult with all stakeholders identified by the department
17 during the rulemaking process. The regulations shall, at a minimum, address all of the
18 following in a manner consistent with this chapter:

19 (A) The process for approving, denying, and disapproving individuals or
20 entities seeking access to information in CURES.

21 (B) The purposes for which a health care practitioner may access information in
22 CURES.

23 (C) The conditions under which a warrant, subpoena, or court order is required
24 for a law enforcement agency to obtain information from CURES as part of a
25 criminal investigation.

26 (D) The process by which information in CURES may be provided for
27 educational, peer review, statistical, or research purposes.

28 (4) In accordance with federal and state privacy laws and regulations, a health
care practitioner may provide a patient with a copy of the patient's CURES patient
activity report as long as no additional CURES data is provided and keep a copy of
the report in the patient's medical record in compliance with subdivision (d) of
Section 11165.1.

(d) For each prescription for a Schedule II, Schedule III, or Schedule IV
controlled substance, as defined in the controlled substances schedules in federal law
and regulations, specifically Sections 1308.12, 1308.13, and 1308.14, respectively, of

Title 21 of the Code of Federal Regulations, the dispensing pharmacy, clinic, or other
dispenser shall report the following information to the Department of Justice as soon
as reasonably possible, but not more than seven days after the date a controlled
substance is dispensed, in a format specified by the Department of Justice:

(1) Full name, address, and, if available, telephone number of the ultimate user
or research subject, or contact information as determined by the Secretary of the
United States Department of Health and Human Services, and the gender, and date of
birth of the ultimate user.

(2) The prescriber's category of licensure, license number, national provider

1 identifier (NPI) number, if applicable, the federal controlled substance registration
2 number, and the state medical license number of any prescriber using the federal
3 controlled substance registration number of a government-exempt facility, if
4 provided.

5 (3) Pharmacy prescription number, license number, NPI number, and federal
6 controlled substance registration number.

7 (4) National Drug Code (N.D.C.) number of the controlled substance dispensed.

8 (5) Quantity of the controlled substance dispensed.

9 (6) International Statistical Classification of Diseases, 9th revision (ICD-9) or
10 10th revision (ICD-10) Code, if available.

11 (7) Number of refills ordered.

12 (8) Whether the drug was dispensed as a refill of a prescription or as a first-time
13 request.

14 (9) Date of origin of the prescription.

15 (10) Date of dispensing of the prescription.

16 (11) The serial number for the corresponding prescription form, if applicable.

17 (e) The Department of Justice may invite stakeholders to assist, advise, and
18 make recommendations on the establishment of rules and regulations necessary to
19 ensure the proper administration and enforcement of the CURES database. All
20 prescriber and dispenser invitees shall be licensed by one of the boards or committees
21 identified in subdivision (d) of Section 208 of the Business and Professions Code, in
22 active practice in California, and a regular user of CURES.

23 (f) The Department of Justice shall, prior to upgrading CURES, consult with
24 prescribers licensed by one of the boards or committees identified in subdivision (d)
25 of Section 208 of the Business and Professions Code, one or more of the boards or
26 committees identified in subdivision (d) of Section 208 of the Business and
27 Professions Code, and any other stakeholder identified by the department, for the
28 purpose of identifying desirable capabilities and upgrades to the CURES Prescription
Drug Monitoring Program (PDMP).

(g) The Department of Justice may establish a process to educate authorized
subscribers of the CURES PDMP on how to access and use the CURES PDMP.

(h) (1) The Department of Justice may enter into an agreement with any entity
operating an interstate data sharing hub, or any agency operating a prescription drug
monitoring program in another state, for purposes of interstate data sharing of
prescription drug monitoring program information.

(2) Data obtained from CURES may be provided to authorized users of another
state's prescription drug monitoring program, as determined by the Department of
Justice pursuant to subdivision (c), if the entity operating the interstate data sharing
hub, and the prescription drug monitoring program of that state, as applicable, have
entered into an agreement with the Department of Justice for interstate data sharing of
prescription drug monitoring program information.

1 (3) Any agreement entered into by the Department of Justice for purposes of
interstate data sharing of prescription drug monitoring program information shall
2 ensure that all access to data obtained from CURES and the handling of data
contained within CURES comply with California law, including regulations, and
3 meet the same patient privacy, audit, and data security standards employed and
required for direct access to CURES.

4 (4) For purposes of interstate data sharing of CURES information pursuant to
this subdivision, and authorized user of another state's prescription drug monitoring
5 program shall not be required to register with CURES, if he or she is registered and in
good standing with that state's prescription drug monitoring program.
6

7 (5) The Department of Justice shall not enter into an agreement pursuant to this
subdivision until the department has issued final regulations regarding the access and
8 use of the information within CURES as required by paragraph (3) of subdivision (c).

9 (i) This section shall remain in effect only until January 1, 2021, and as of that
date is repealed.

10 13. Health and Safety Code section 11165.1 states:

11 (a) (1) (A) (i) A health care practitioner authorized to prescribe, order,
administer, furnish, or dispense Schedule II, Schedule III, or Schedule IV controlled
12 substances pursuant to Section 11150 shall, before July 1, 2016, or upon receipt of a
federal Drug Enforcement Administration (D.E.A.) registration, whichever occurs
13 later, submit an application developed by the Department of Justice to obtain
approval to electronically access information regarding the controlled substance
14 history of a patient that is maintained by the Department of Justice. Upon approval,
the department shall release to that practitioner the electronic history of controlled
15 substances dispensed to an individual under the practitioner's care based on data
contained in the CURES Prescription Drug Monitoring Program (PDMP).
16

17 (ii) A pharmacist shall, before July 1, 2016, or upon licensure, whichever
occurs later, submit an application developed by the Department of Justice to obtain
18 approval to electronically access information regarding the controlled substance
history of a patient that is maintained by the Department of Justice. Upon approval,
19 the department shall release to that pharmacist the electronic history of controlled
substances dispensed to an individual under the practitioner's care based on data
20 contained in the CURES PDMP.

21 (B) An application may be denied, or a subscriber may be suspended, for
reasons which include, but are not limited to, the following:

22 (i) Materially falsifying an application to access information contained in the
CURES database.
23

24 (ii) Failing to maintain effective controls for access to the patient activity
report.

25 (iii) Having his or her federal D.E.A. registration suspended or revoked.

26 (iv) Violating a law governing controlled substances or any other law for which
the possession or use of a controlled substance is an element of the crime.
27

28 (v) Accessing information for a reason other than to diagnose or treat his or her
patients, or to document compliance with the law.

1 (C) An authorized subscriber shall notify the Department of Justice within 30
2 days of any changes to the subscriber account.

3 (D) Commencing no later than October 1, 2018, an approved health care
4 practitioner, pharmacist, and any person acting on behalf of a health care practitioner
5 or pharmacist pursuant to subdivision (b) of Section 209 of the Business and
6 Professions Code may use the department's online portal or a health information
7 technology system that meets the criteria required in subparagraph (E) to access
8 information in the CURES database pursuant to this section. A subscriber who uses a
9 health information technology system that meets the criteria required in subparagraph
10 (E) to access the CURES database may submit automated queries to the CURES
11 database that are triggered by predetermined criteria.

12 (E) Commencing no later than October 1, 2018, an approved health care
13 practitioner or pharmacist may submit queries to the CURES database through a
14 health information technology system if the entity that operates the health information
15 technology system can certify all of the following:

16 (i) The entity will not use or disclose data received from the CURES database
17 for any purpose other than delivering the data to an approved health care practitioner
18 or pharmacist or performing data processing activities that may be necessary to
19 enable the delivery unless authorized by, and pursuant to, state and federal privacy
20 and security laws and regulations.

21 (ii) The health information technology system will authenticate the identity of
22 an authorized health care practitioner or pharmacist initiating queries to the CURES
23 database and, at the time of the query to the CURES database, the health information
24 technology system submits the following data regarding the query to CURES:

25 (I) The date of the query.

26 (II) The time of the query.

27 (III) The first and last name of the patient queried.

28 (IV) The date of birth of the patient queried.

(V) The identification of the CURES user for whom the system is making the
query.

(iii) The health information technology system meets applicable patient privacy
and information security requirements of state and federal law.

(iv) The entity has entered into a memorandum of understanding with the
department that solely addresses the technical specifications of the health information
technology system to ensure the security of the data in the CURES database and the
secure transfer of data from the CURES database. The technical specification shall
be universal for all health information technology systems that establish a method of
system integration to retrieve information from the CURES database. The
memorandum of understanding shall not govern, or in any way impact or restrict, the
use of data received from the CURES database or impose any additional burdens on
covered entities in compliance with regulations promulgated pursuant to the federal
Health Insurance Portability and Accountability Act of 1996 found in Parts 160 and
164 of Title 45 of the Code of Federal Regulations.

(F) No later than October 1, 2018, the department shall develop a programming

1 interface or other method of system integration to allow health information
2 technology systems that meet the requirements in subparagraph (E) to retrieve
3 information in the CURES database on behalf of an authorized health care
4 practitioner or pharmacist.

5 (G) The department shall not access patient-identifiable information in an
6 entity's health information technology system.

7 (H) An entity that operates a health information technology system that is
8 requesting to establish an integration with the CURES database shall pay a reasonable
9 fee to cover the costs of establishing and maintaining integration with the CURES
10 database.

11 (I) The department may prohibit integration or terminate a health information
12 technology system's ability to retrieve information in the CURES database if the
13 health information technology system fails to meet the requirements of subparagraph
14 (E), or the entity operating the health information technology system does not fulfill
15 its obligation under subparagraph (H).

16 (2) A health care practitioner authorized to prescribe, order, administer, furnish,
17 or dispense Schedule II, Schedule III, or Schedule IV controlled substances pursuant
18 to Section 11150 or a pharmacist shall be deemed to have complied with paragraph
19 (1) if the licensed health care practitioner or pharmacist has been approved to access
20 the CURES database through the process developed pursuant to subdivision (a) of
21 Section 209 of the Business and Professions Code.

22 (b) A request for, or release of, a controlled substance history pursuant to this
23 section shall be made in accordance with guidelines developed by the Department of
24 Justice.

25 (c) In order to prevent the inappropriate, improper, or illegal use of Schedule II,
26 Schedule III, or Schedule IV controlled substances, the Department of Justice may
27 initiate the referral of the history of controlled substances dispensed to an individual
28 based on data contained in CURES to licensed health care practitioners, pharmacists,
or both, providing care or services to the individual.

(d) The history of controlled substances dispensed to an individual based on
data contained in CURES that is received by a practitioner or pharmacist from the
Department of Justice pursuant to this section is medical information subject to the
provisions of the Confidentiality of Medical Information Act contained in Part 2.6
(commencing with Section 56) of Division 1 of the Civil Code.

(e) Information concerning a patient's controlled substance history provided to
a prescriber or pharmacist pursuant to this section shall include prescriptions for
controlled substances listed in Sections 1308.12, 1308.13, and 1308.14 of Title 21 of
the Code of Federal Regulations.

(f) A health care practitioner, pharmacist, and any person acting on behalf of a
health care practitioner or pharmacist, when acting with reasonable care and in good
faith, is not subject to civil or administrative liability arising from any false,
incomplete, inaccurate, or misattributed information submitted to, reported by, or
relied upon in the CURES database or for any resulting failure of the CURES
database to accurately or timely report that information.

(g) For purposes of this sections, the following terms have the following
meanings:

1 (1) "Automated basis" means using predefined criteria to trigger an automated
2 query to the CURES database, which can be attributed to a specific health care
3 practitioner or pharmacist.

4 (2) "Department" means the Department of Justice.

5 (3) "Entity" means an organization that operates, or provides or makes
6 available, a health information technology system to health care practitioner or
7 pharmacist.

8 (4) "Health information technology system" means an information processing
9 application using hardware and software for the storage, retrieval, sharing of or use of
10 patient data for communication, decision making, coordination of care, or the quality,
11 safety, or efficiency of the practice of medicine or delivery of health care services,
12 including, but not limited to, electronic medical record applications, health
13 information exchange systems, or other interoperable clinical or health care
14 information system.

15 (5) "User initiated basis" means an authorized health care practitioner or
16 pharmacist has taken an action to initiate the query to the CURES database, such as
17 clicking a button, issuing a voice command, or taking some other action that can be
18 attributed to a specific health care practitioner or pharmacist.

19 (h) This section shall become inoperative on July 1, 2021, or upon the date the
20 department promulgates regulations to implement this section and posts those
21 regulations on its internet website, whichever date is earlier, and, as of January 1,
22 2022, is repealed.

23 14. Health and Safety Code section 11165.4 states:

24 (a) (1) (A) (i) A health care practitioner authorized to prescribe, order,
25 administer, or furnish a controlled substance shall consult the CURES database to
26 review a patient's-controlled substance history before prescribing a Schedule II,
27 Schedule III, or Schedule IV controlled substance to the patient for the first time and
28 at least once every four months thereafter if the substance remains part of the
treatment of the patient.

(ii) If a health care practitioner authorized to prescribe, order, administer, or
furnish a controlled substance is not required, pursuant to an exemption described in
subdivision (c), to consult the CURES database the first time he or she prescribes,
orders, administers, or furnishes a controlled substance to a patient, he or she shall
consult the CURES database to review the patient's controlled substance history
before subsequently prescribing a Schedule II, Schedule III, or Schedule IV
controlled substance to the patient and at least once every four months thereafter if
the substance remains part of the treatment of the patient.

(B) For purposes of this paragraph, first time means the initial occurrence in
which a health care practitioner, in his or her role as a health care practitioner, intends
to prescribe, order, administer, or furnish a Schedule II, Schedule III, or Schedule IV
controlled substance to a patient and has not previously prescribed a controlled
substance to the patient.

(2) A health care practitioner shall obtain a patient's controlled substance
history from the CURES database no earlier than 24 hours, or the previous business
day, before he or she prescribes, orders, administers, or furnishes a Schedule II,
Schedule III, or Schedule IV controlled substance to the patient.

1 (b) The duty to consult the CURES database, as described in subdivision (a),
2 does not apply to veterinarians or pharmacists.

3 (c) The duty to consult the CURES database, as described in subdivision (a),
4 does not apply to a health care practitioner in any of the following circumstances:

5 (1) If a health care practitioner prescribes, orders, or furnishes a controlled
6 substance to be administered to a patient while the patient is admitted to any of the
7 following facilities or during an emergency transfer between any of the following
8 facilities for use while on facility premises:

9 (A) A licensed clinic, as described in Chapter 1 (commencing with Section
10 1200) of Division 2.

11 (B) An outpatient setting, as described in Chapter 1.3 (commencing with
12 Section 1248) of Division 2.

13 (C) A health facility, as described in Chapter 2 (commencing with Section
14 1250) of Division 2.

15 (D) A county medical facility, as described in Chapter 2.5 (commencing with
16 Section 1440) of Division 2.

17 (2) If a health care practitioner prescribes, orders, administers, or furnishes a
18 controlled substance in the emergency department of a general acute care hospital and
19 the quantity of the controlled substance does not exceed a nonrefillable seven-day
20 supply of the controlled substance to be used in accordance with the directions for
21 use.

22 (3) If a health care practitioner prescribes, orders, administers, or furnishes a
23 controlled substance to a patient as part of the patient's treatment for a surgical
24 procedure and the quantity of the controlled substance does not exceed a nonrefillable
25 five-day supply of the controlled substance to be used in accordance with the
26 directions for use, in any of the following facilities:

27 (A) A licensed clinic, as described in Chapter 1 (commencing with Section
28 1200) of Division 2.

(B) An outpatient setting, as described in Chapter 1.3 (commencing with
Section 1248) of Division 2.

(C) A health facility, as described in Chapter 2 (commencing with Section
1250) of Division 2.

(D) A county medical facility, as described in Chapter 2.5 (commencing with
Section 1440) of Division 2.

(E) A place of practice, as defined in Section 1658 of the Business and
Professions Code.

(4) If a health care practitioner prescribes, orders, administers, or furnishes a
controlled substance to a patient currently receiving hospice care, as defined in
Section 1339.40.

(5) (A) If all of the following circumstances are satisfied:

1 (i) It is not reasonably possible for a health care practitioner to access the
information in the CURES database in a timely manner.

2 (ii) Another health care practitioner or designee authorized to access the
3 CURES database is not reasonably available.

4 (iii) The quantity of controlled substance prescribed, ordered, administered, or
5 furnished does not exceed a nonrefillable five-day supply of the controlled substance
to be used in accordance with the directions for use and no refill of the controlled
substance is allowed.

6 (B) A health care practitioner who does not consult the CURES database under
7 subparagraph (A) shall document the reason he or she did not consult the database in
the patient's medical record.

8 (6) If the CURES database is not operational, as determined by the department,
9 or when it cannot be accessed by a health care practitioner because of a temporary
technological or electrical failure. A health care practitioner shall, without undue
10 delay, seek to correct any cause of the temporary technological or electrical failure
that is reasonably within his or her control.

11 (7) If the CURES database cannot be accessed because of technological
12 limitations that are not reasonably within the control of a health care practitioner.

13 (8) If consultation of the CURES database would, as determined by the health
14 care practitioner, result in a patient's inability to obtain a prescription in a timely
manner and thereby adversely impact the patient's medical condition, provided that
15 the quantity of the controlled substance does not exceed a nonrefillable five-day
supply if the controlled substance were used in accordance with the directions for use.

16 (d) (1) A health care practitioner who fails to consult the CURES database, as
17 described in subdivision (a), shall be referred to the appropriate state professional
licensing board solely for administrative sanctions, as deemed appropriate by that
board.

18 (2) This section does not create a private cause of action against a health care
19 practitioner. This section does not limit a health care practitioner's liability for the
negligent failure to diagnose or treat a patient.

20 (e) This section is not operative until six months after the Department of Justice
21 certifies that the CURES database is ready for statewide use and that the department
has adequate staff, which, at a minimum, shall be consistent with the appropriation
22 authorized in Schedule (6) of Item 0820-001-0001 of the Budget Act of 2016
(Chapter 23 of the Statutes of 2016), user support, and education. The department
23 shall notify the Secretary of State and the office of the Legislative Counsel of the date
of that certification.

24 (f) All applicable state and federal privacy laws govern the duties required by
25 this section.

26 (g) The provisions of this section are severable. If any provision of this section
or its application is held invalid, that invalidity shall not affect other provisions or
27 applications that can be given effect without the invalid provision or application.

28 (h) This section shall become inoperative on July 1, 2021, or upon the date the
department promulgates regulations to implement this section and posts those

1 regulations on its internet website, whichever date is earlier, and, as of January 1,
2 2022, is repealed.

3 15. Section 2228.1 of the Code provides as follows:

4 (a) On and after July 1, 2019, except as otherwise provided in subdivision (c),
5 the board shall require a licensee to provide a separate disclosure that includes the
6 licensee's probation status, the length of the probation, the probation end date, all
7 practice restrictions placed on the licensee by the board, the board's telephone
8 number, and an explanation of how the patient can find further information on the
9 licensee's probation on the licensee's profile page on the board's online license
10 information Internet Web site, to a patient or the patient's guardian or health care
11 surrogate before the patient's first visit following the probationary order while the
12 licensee is on probation pursuant to a probationary order made on and after July 1,
13 2019, in any of the following circumstances:

14 (1) A final adjudication by the board following an administrative hearing or
15 admitted findings or prima facie showing in a stipulated settlement establishing any
16 of the following:

17 (A) The commission of any act of sexual abuse, misconduct, or relations with a
18 patient or client as defined in Section 726 or 729.

19 (B) Drug or alcohol abuse directly resulting in harm to patients or the extent
20 that such use impairs the ability of the licensee to practice safely.

21 (C) Criminal conviction directly involving harm to patient health.

22 (D) Inappropriate prescribing resulting in harm to patients and a probationary
23 period of five years or more.

24 (2) An accusation or statement of issues alleged that the licensee committed any
25 of the acts described in subparagraphs (A) to (D), inclusive, of paragraph (1), and a
26 stipulated settlement based upon a nolo contendere or other similar compromise that
27 does not include any prima facie showing or admission of guilt or fact but does
28 include an express acknowledgment that the disclosure requirements of this section
would serve to protect the public interest.

(b) A licensee required to provide a disclosure pursuant to subdivision (a) shall
obtain from the patient, or the patient's guardian or health care surrogate, a separate,
signed copy of that disclosure.

(c) A licensee shall not be required to provide a disclosure pursuant to
subdivision (a) if any of the following applies:

(1) The patient is unconscious or otherwise unable to comprehend the
disclosure and sign the copy of the disclosure pursuant to subdivision (b) and a
guardian or health care surrogate is unavailable to comprehend the disclosure and
sign the copy.

(2) The visit occurs in an emergency room or an urgent care facility or the visit
is unscheduled, including consultations in inpatient facilities.

(3) The licensee who will be treating the patient during the visit is not known to
the patient until immediately prior to the start of the visit.

1 (4) The licensee does not have a direct treatment relationship with the patient.

2 (d) On and after July 1, 2019, the board shall provide the following
3 information, with respect to licensees on probation and licensees practicing under
4 probationary licenses, in plain view on the licensee's profile page on the board's
5 online license information Internet Web site.

6 (1) For probation imposed pursuant to a stipulated settlement, the causes
7 alleged in the operative accusation along with a designation identifying those causes
8 by which the licensee has expressly admitted guilt and a statement that acceptance of
9 the settlement is not an admission of guilt.

10 (2) For probation imposed by an adjudicated decision of the board, the causes
11 for probation stated in the final probationary order.

12 (3) For a licensee granted a probationary license, the causes by which the
13 probationary license was imposed.

14 (4) The length of the probation and end date.

15 (5) All practice restrictions placed on the license by the board.

16 (e) Section 2314 shall not apply to this section.

17 FACTUAL ALLEGATIONS

18 PATIENT 1

19 16. An inquiry into the care of Patient 1 commenced after a complaint was submitted to
20 the Board by the daughter of Patient 1¹, which alleged that Respondent had been prescribing
21 medications to her father for a period of more than two years even though Patient 1 was already
22 under the care of an assisted living facility for his medical conditions and medication
23 management. The daughter of Patient 1, who held a health care power of attorney for her father,
24 stated that she had asked Respondent on numerous occasions to stop prescribing multiple
25 medications to her father. However, Respondent had refused to stop prescribing medications to
26 Patient 1, citing "social ties" as the reason to continue prescribing. This conduct, the daughter
27 alleged, caused her father to have an increasing number of falling incidents. Additionally, the
28 daughter alleged that her father's medical conditions could not be effectively managed while
Respondent continued to prescribe medications. Patient 1 was prescribed the following

¹ The individual patients are referred to by numbers to protect patient privacy. Their identity will be disclosed to the Respondent in discovery.

1 medications by Respondent: Xanax, Dexedrine, Abilify, Ambien, nitroglycerin, opioids, and
2 medication for diabetes (a condition which Patient 1 did not have).

3 17. The records provided by Respondent for Patient 1 total approximately 47 pages from
4 the period beginning approximately October 15, 2015 through approximately March 5, 2018.
5 Although Patient 1 had been a patient of Respondent's since approximately April 4, 2012,
6 Respondent was unable to provide records for the dates between approximately April 4, 2012
7 through approximately October 15, 2015, as a result of the records having been destroyed. As
8 such, there are no clinic notes or visits for the time period between approximately April 4, 2012
9 through approximately October 15, 2015. However, the CURES System recorded that controlled
10 substances were prescribed to Patient 1 by Respondent from approximately April 9, 2012, to
11 approximately March 6, 2018. According to CURES, Patient 1 was routinely prescribed by
12 Respondent the following medications: Alprazolam 1 mg., Zolpidem Tartrate 10 mg.,
13 Dextroamphetamine Sulfate 15 mg., Tramadol 50 mg., Temazepam 30 mg., and Testosterone
14 Cypionate 200 mg. According to records from the nursing home and other outside hospitals,
15 Patient 1 was a 78-year-old male with a history of coronary artery disease, schizophrenia, anxiety,
16 hypertension, hyperlipidemia, benign prostatic hypertrophy, multiple falls, and pressure ulcers.

17 18. The medical records from Respondent's office reveal illegible progress notes with
18 minimal documentation regarding Patient 1's medical condition. For instance, it is impossible to
19 ascertain the chief complaint or reason for the medical visit in the progress notes. The section of
20 the progress note for current medications was left blank. Vital signs were not consistently
21 documented. The physical examination portion of the progress notes is illegible, making it
22 impossible to determine what was examined on the date of service. Lastly, the assessment and
23 diagnosis sections were illegible, making it impossible to determine what was prescribed to the
24 patient and for what specific indication. The documentation lacked any pain assessment, lacked
25 any assessment to determine if there was a prior substance abuse history, and there was no
26 indication in the medical records that Respondent ever evaluated a CURES report to determine if
27 Patient 1 was being prescribed controlled substances by another physician. Further, a clear
28 indication for prescribing the controlled substances was not documented. There was no evidence

1 in the medical records that Respondent had made any referrals or diagnostic studies related to
2 prescribing the controlled substances for this patient.

3 19. The standard of care requires a medical history and physical exam, which includes an
4 assessment of the patient's pain, including physical and psychological status and function;
5 substance abuse history; history of prior pain treatments and assessment of any other underlying
6 or coexisting conditions. Finally, it should include documentation of recognized medical
7 indications for the use of controlled substances. A history and physical was not documented in
8 any clinic visits by Respondent. The period between approximately April 4, 2012 through
9 approximately October 15, 2015, had no corresponding medical records, although the CURES
10 report for that time period noted that Respondent had prescribed or refilled approximately 62
11 prescriptions for controlled substances. There was no evidence that a substance abuse history was
12 performed in the medical records. The records were consistently illegible and lacked
13 documentation of clearly recognized medical indications for the use of controlled substances
14 given the patient's medical conditions. The records fail to document a review of systems and the
15 lack of a physical examination. There was no clear indication for Respondent's prescribing of
16 controlled substances to treat Patient 1's known medical conditions.

17 20. The lack of a documented substance abuse history and lack of a review of systems
18 and physical examination constitute an extreme departure from the standard of care. The
19 prescribing of controlled substances without a clear medical indication for the controlled
20 substance is also an extreme departure from the standard of care. The lack of medical records for
21 an approximately three-and-a-half-year period during which a patient was prescribed
22 approximately 62 controlled substances is an extreme departure from the standard of care.

23 21. The standard of care requires the medical records contain stated objectives that may
24 include relief of pain or relief of the medical condition requiring controlled substances and/or
25 improved physical or psychological function or ability to perform certain tasks or activities of
26 daily living. This should also include any plans for further diagnostic evaluations and treatments,
27 such as a rehabilitation program. The medical records for the period of approximately October 15,
28 2015 through approximately March 5, 2018, reflect repeated departures from the standard of care

1 given Respondent's failure to document a specific treatment plan for this patient. The medical
2 records did not document any treatment plans and had no clear plan or objectives for the ongoing
3 prescribing of controlled substances.

4 22. Respondent's failure to document a specific treatment plan for this patient constitutes
5 repeated extreme departures from the standard of care. There was no evidence that Respondent
6 ordered any additional diagnostic evaluations or treatment for the patient related to the indications
7 for prescribing the numerous controlled substances.

8 23. The standard of care requires that the medical records document that the physician
9 discussed the risks and benefits of using controlled substances and other treatment modalities. An
10 actual written consent is not required but is recommended. There was no evidence in the medical
11 records that Respondent discussed potential side effects and risks of controlled substances. There
12 were no written consent forms noted in the medical records. Respondent's clinical notes do not
13 indicate that the risks of controlled substances were discussed. This constitutes an extreme
14 departure from the standard of care as there was no evidence in the medical records that
15 Respondent discussed the potential side effects and risks of ongoing prescribing of controlled
16 substances to Patient 1.

17 24. The standard of care requires the medical records reflect that the physician is
18 periodically reviewing the course of treatment and his prescribing of controlled substances for the
19 patient and making appropriate modifications in treatment based on the patient's progress or lack
20 of progress. However, the medical records for Patient 1 fail to demonstrate that Respondent ever
21 performed a periodic review on the patient's ongoing treatment with controlled substances even
22 though the patient was prescribed multiple controlled substances over a period of approximately
23 seven years.

24 25. Respondent's failure to perform periodic reviews of the patient's treatment and status
25 over a period of approximately seven years in the setting of repeated prescribing and refilling of
26 multiple controlled substances constitutes multiple extreme departures from the standard of care.

27 26. The standard of care requires the physician consider obtaining additional evaluations
28 and consultations, especially with complex pain problems. Special attention should be given to

1 patients who are at risk for misusing their medication or have a history of drug addiction or
2 substance abuse. Such patients require extra care and monitoring and documentation and
3 consultation with an addiction medicine specialist and, if indicated, a pain management specialist.
4 There is no evidence in the medical records that Respondent made any referrals or obtained a
5 consultation related to prescribing the controlled substances for Patient 1. There is no evidence
6 that Respondent had assessed the patient for any prior history of drug addiction or substance
7 abuse. Additionally, there is no evidence that Respondent ever requested or reviewed a CURES
8 report to determine if Patient 1 was being prescribed controlled substances by another physician.

9 27. Respondent's failure to obtain a consultation for Patient 1 was a simple departure
10 from the standard of care. Respondent's failure to determine if the patient was prescribed
11 controlled substances by another provider was an extreme departure from the standard of care.
12 Respondent did not employ CURES to determine if a patient he continuously prescribed
13 controlled substances to for more than approximately seven years was receiving controlled
14 substances from other providers.

15 28. The standard of care requires a physician to maintain accurate and complete records,
16 demonstrate a history and exam, evaluations and consultations, treatment plans and objectives,
17 informed consent, medications prescribed, and periodic review documentation. The medical
18 records for all visits are largely illegible in all sections of the progress note. The review of
19 systems and physical examination sections of the progress notes are also largely illegible and fail
20 to document why the patient was being seen, what part of the physical examination was
21 performed, and the specific assessment and plan of care for the patient. The records lacked
22 medical indication for the medications prescribed over a period of approximately seven years.

23 29. It was an extreme departure from the standard of care to have failed to produce a
24 medical record for Patient 1 that documented standard guidelines in the use of controlled
25 substances.

26 30. Through inappropriate prescribing of controlled substances, Respondent caused harm
27 to Patient 1. The inappropriate prescribing of controlled substances (in this case Xanax,
28 Dexedrine, Ambien, and opioid medication) by Respondent, without proper justification or

1 medical indication for such substances led to placing Patient 1 at an unnecessarily increased risk
2 for significant morbidity and mortality and potential harm given his pre-existing chronic medical
3 conditions. Specifically, the following harm resulted. Patient 1 had a history of hypertension and
4 coronary artery disease. He was prescribed multiple controlled substances that had a high
5 potential for abuse and dependency, which most likely resulted in Patient 1 developing a
6 dependency on multiple controlled substances that were unnecessarily prescribed. Additionally,
7 medication such as Dexedrine can lead to increased blood pressure which could potentially lead
8 to adverse effects in a patient with known hypertension and coronary artery disease. Patient 1
9 was harmed by the Respondent's inappropriate prescribing. Patient 1 unnecessarily developed a
10 likely dependency on multiple controlled substances, which should not have been prescribed to
11 him given the lack of medical justification or medical indication for inappropriately prescribing of
12 such medications by Respondent.

13 PATIENT 2

14 31. Patient 2 was an adult female who had a history of insomnia, depression, anxiety, and
15 back pain. On or about June 28, 2018, the patient was seen by Respondent for complaints of
16 nasal blockage, back and neck pain, dermatitis, a swollen nose, and muscle spasms. From
17 approximately 2016 to approximately 2018, Patient 2 was prescribed consistently the following
18 medications by Respondent: Hydrocodone 10 mg., Alprazolam 1mg., and Zolpidem 10 mg.

19 32. Respondent failed to comply with the standard of care for prescribing controlled
20 substances to Patient 2, who had a history of insomnia, depression, anxiety, and back pain. The
21 standard of care under such circumstances requires a medical history and physical exam, which
22 includes an assessment of the patient's pain, including physical and psychological status and
23 function; substance abuse history; history of prior pain treatments and assessment of any other
24 underlying or co-existing conditions. Finally, it should include documentation of recognized
25 medical indications for the use of controlled substances.

26 33. A history and physical was not documented in any clinic visits by Respondent. A
27 review of the Respondent's medical records for Patient 2 for the period between approximately
28 February 1, 2016 through approximately June 30, 2018, provide no evidence that a substance

1 abuse history was performed. The records are consistently illegible and lack documentation of
2 clearly recognized medical indications for the use of controlled substances given the patient's
3 medical conditions. In addition, the medical records of Patient 2 show a lack of a review of
4 systems and the lack of a suitable physical examination. There was no clear indication
5 documented for Respondent's prescribing of controlled substances for treatment of Patient 2's
6 known medical conditions.

7 34. The lack of a documented substance abuse history and lack of a review of systems
8 and physical examination constitute an extreme departure from the standard of care. The
9 prescribing of controlled substances without a clear medical indication for the controlled
10 substance is an extreme departure from the standard of care.

11 35. The standard of care requires the medical records contain stated objectives that may
12 include relief of pain or relief of the medical condition requiring controlled substances and/or
13 improved physical or psychological function or ability to perform certain tasks or daily living
14 activities. This should also include any plans for further diagnostic evaluations and treatments,
15 such as a rehabilitation program.

16 36. From approximately February 1, 2016 through approximately June 30, 2018, the
17 medical records fail to document a specific treatment plan for this patient. The medical records
18 do not document any treatment plans or any clear plan or objectives for the ongoing prescribing
19 of controlled substances.

20 37. Respondent's failure to document a specific treatment plan for this patient was an
21 extreme departure from the standard of care. There was no evidence documented in the medical
22 record for Patient 2 to support that Respondent ordered any additional diagnostic evaluations or
23 treatment for the patient related to the indications for prescribing the numerous controlled
24 substances.

25 38. The standard of care requires the medical records document that the physician discuss
26 risks and benefits of using controlled substances and other treatment modalities. An actual
27 written consent is not required but is recommended. There was no evidence in the medical
28 records that Respondent discussed potential side effects and risks of controlled substances. There

1 were no written consent forms noted in the medical records. Respondent's clinical notes do not
2 indicate that the risks of controlled substances were discussed. This is an extreme departure from
3 the standard of care as there was no evidence in the medical records that Respondent discussed
4 the potential side effects and risks of ongoing prescribing of controlled substances to Patient 2.

5 39. The standard of care requires that patient records reflect that the physician is
6 periodically reviewing the course of treatment and his prescribing of controlled substances for the
7 patient, and making appropriate modifications in treatment based on the patient's progress or lack
8 of progress. The medical records on Patient 2 do not demonstrate that Respondent ever
9 performed a periodic review on the patient's ongoing treatment with controlled substances even
10 though the patient was prescribed multiple controlled substances over a period of approximately
11 two years.

12 40. Respondent's failure over a period of approximately two years to perform periodic
13 reviews of the patient's treatment and status in the setting of repeated prescribing and refilling of
14 multiple controlled substances constitutes multiple extreme departures from the standard of care.

15 41. The standard of care requires that the physician consider obtaining additional
16 evaluations and consultations, especially with complex pain problems. Special attention should
17 be given to patients who are at risk for misusing their medication or have a history of drug
18 addiction or substance abuse. Such patients require extra care and monitoring and
19 documentation, and consultation with an addiction medicine specialist and, if indicated, a pain
20 management specialist.

21 42. There is no evidence in the medical records that Respondent had made any referrals
22 or obtained a consultation related to prescribing the controlled substances for this patient. There
23 is no evidence that Respondent had assessed the patient for any prior history of drug addiction or
24 substance abuse.

25 43. Respondent's failure to obtain a consultation related to prescribing controlled
26 substances for Patient 2 was a simple departure from the standard of care.

27 44. The standard of care requires the physician must maintain accurate and complete
28 records, demonstrating a history and exam along with evaluations and consultations, treatment

1 plans and objectives, informed consent, medications prescribed, and periodic review
2 documentation.

3 45. The medical records and progress notes for all visits with Patient 2 are largely
4 illegible in all sections of the progress note. Also, the review of systems and physical
5 examination sections of the progress notes are illegible, so that a reader could not determine why
6 the patient was being seen, what part of the physical examination was performed and what the
7 specific assessment and plan of care was for the patient. In addition, the records lack medical
8 indication for the medications prescribed over a period of approximately two years.

9 46. Respondent's failure to document standard guidelines in the use of controlled
10 substances for Patient 2 was an extreme departure from the standard of care.

11 47. Through inappropriate prescribing of controlled substances, Respondent harmed
12 Patient 2. The inappropriate prescribing of controlled substances (Hydrocodone 10 mg.,
13 Alprazolam 1mg., and Zolpidem 10 mg.) by Respondent, without proper justification or medical
14 indication for such substances over a period of approximately three years, led to placing Patient 2,
15 a patient with a history of depression, anxiety, and back pain at an unnecessarily increased risk
16 for significant morbidity and mortality and potential harm, including accelerated progression of
17 her pre-existing chronic medical conditions and ongoing dependency on controlled substances.
18 Specific harm resulted from the Respondent's conduct. Patient 2 had a history of depression,
19 anxiety, and back pain. She was prescribed multiple controlled substances by Respondent that
20 had a high potential for abuse and dependency in a patient with multiple medical conditions who
21 was at risk for exacerbation of said co-morbidities by taking unnecessarily prescribed controlled
22 medications. This resulted in Patient 2 developing a dependency on multiple controlled
23 substances that were prescribed without medical indication. The Respondent's inappropriate
24 prescribing resulted in patient harm to Patient 2. Patient 2 unnecessarily developed a dependency
25 on multiple controlled substances which should not have been prescribed to her given the lack of
26 medical justification or medical indication for inappropriately prescribing of such medications by
27 Respondent.

28 //

1 PATIENT 3

2 48. Patient 3 was an adult male patient with a history of insomnia, anxiety, dizziness,
3 dermatitis, cellulitis, back pain, hypertension, hearing loss, depression, and benign prostate
4 hypertrophy. The Respondent prescribed the following medications to Patient 3 on a consistent
5 basis during the period from approximately 2015 to approximately 2018: clonazepam,
6 hydrocodone, zolpidem, and Soma.

7 49. Under such circumstances, the standard of care requires a medical history and
8 physical exam, which includes an assessment of the patient's pain, including physical and
9 psychological status and function; substance abuse history; history of prior pain treatments and
10 assessment of any other underlying or co-existing conditions. Finally, it should include
11 documentation of recognized medical indications for the use of controlled substances. A history
12 and physical was not documented in any clinic visits by Respondent. There was no evidence that
13 a substance abuse history was performed in the medical records between approximately July 1,
14 2015, through approximately December 31, 2018. The records were consistently illegible and
15 lacked documentation of clearly recognized medical indications for the use of controlled
16 substances given the patient's medical conditions. The records showed a lack of a review of
17 systems and the lack of a physical examination. There was no clear indication for Respondent's
18 prescribing of controlled substances for treatment of Patient 3's known medical conditions.

19 50. The lack of a documented substance abuse history and lack of a review of systems
20 and physical examination constitute an extreme departure from the standard of care. The
21 prescribing of controlled substances without a clear medical indication for the controlled
22 substance is an extreme departure from the standard of care.

23 51. The standard of care requires the medical records contain stated objectives that may
24 include relief of pain or relief of the medical condition requiring controlled substances and/or
25 improved physical or psychological function or ability to perform certain tasks or activities of
26 daily living. This should also include any plans for further diagnostic evaluations and treatments,
27 such as a rehabilitation program. The medical records from the period of approximately July 1,
28 2015 through approximately December 31, 2018, reflect repeated departures from the standard of

1 care given Respondent's failure to document a specific treatment plan for this patient. The
2 medical records do not have a clear plan or objectives for the ongoing prescribing of controlled
3 substances.

4 52. Respondent's repeated failure to document a specific treatment plan for this patient
5 constitutes extreme departures from the standard of care. There was no evidence to support that
6 Respondent ordered any additional diagnostic evaluations or treatment for the patient related to
7 the indications for prescribing the numerous controlled substances.

8 53. The standard of care requires the medical records document that the physician
9 discussed risks and benefits of the use of controlled substances along with other treatment
10 modalities. An actual written consent is not required but is recommended. There was no
11 evidence in the medical records that Respondent discussed potential side effects and risks of
12 controlled substances. There were no written consent forms in the medical records.

13 54. Respondent's clinical notes do not indicate that the risks of controlled substances
14 were discussed. This constitutes an extreme departure from the standard of care as there was no
15 evidence in the medical records that Respondent discussed the potential side effects and risks of
16 ongoing prescribing of controlled substances to Patient 3.

17 55. The standard of care requires the medical records reflect that the physician
18 periodically review the course of treatment and prescribing of controlled substances for the
19 patient and making appropriate modifications in treatment based on the patient's progress or lack
20 of progress. The medical records on Patient 3 fail to demonstrate that Respondent ever performed
21 a periodic review on the patient's ongoing treatment with controlled substances even though the
22 patient was prescribed multiple controlled substances over a period of over approximately three
23 years.

24 56. There were multiple extreme departures from the standard of care over a period of
25 over approximately three years for failure to perform periodic reviews of the patient's treatment
26 and status in the setting of repeated prescribing and refilling of multiple controlled substances.

27 57. The standard of care under such circumstances requires that a physician consider
28 obtaining additional evaluations and consultations, especially with complex pain problems.

1 Special attention should be given to patients who are at risk for misusing their medication or have
2 a history of drug addiction or substance abuse. Such patients require extra care and monitoring
3 and documentation, and consultation with an addiction medicine specialist and, if indicated, a
4 pain management specialist. There is no evidence in the medical records for Patient 3 that
5 Respondent made any referrals or obtained a consultation related to prescribing the controlled
6 substances for this patient. There is no evidence that Respondent had assessed the patient for any
7 prior history of drug addiction or substance abuse.

8 58. Respondent's failure to obtain a consultation in this patient is a simple departure from
9 the standard of care.

10 59. The standard of care requires the physician to maintain accurate and complete
11 records, demonstrate a history and exam, evaluations and consultations, treatment plans and
12 objectives, informed consent, medications prescribed, and periodic review documentation. The
13 medical records for all visits with Patient 3 are largely illegible in all sections of the progress
14 note. The review of systems and physical examination sections of the progress notes are likewise
15 largely illegible, making it impossible to determine why the patient was being seen, what part of
16 the physical examination was performed and what the specific assessment and plan of care was
17 for the patient. The records lacked medical indication for the medications prescribed over a
18 period of over approximately three years.

19 60. Respondent's failure to document standard guidelines in the use of controlled
20 substances for Patient 3 is an extreme departure from the standard of care.

21 61. In addition, the inappropriate prescribing of controlled substances (clonazepam,
22 hydrocodone, zolpidem, and Soma) by Respondent to Patient 3, without adequate justification or
23 medical indication for such substances over a period of approximately three years, harmed Patient
24 3 by placing him at an unnecessarily increased risk for significant morbidity and mortality and
25 potential harm given his pre-existing chronic medical conditions. Patient 3 had a history of
26 insomnia, anxiety, dizziness, dermatitis, cellulitis, back pain, hypertension, hearing loss,
27 depression, and benign prostate hypertrophy. He was prescribed multiple controlled substances
28 by Respondent that had high potential for abuse and dependency in a patient with multiple

1 medical conditions who was at risk for exacerbation of said co-morbidities by taking
2 unnecessarily prescribed controlled medications. This resulted in Patient 3 developing a
3 dependency on multiple controlled substances that were prescribed without medical indication.
4 Patient 3 unnecessarily developed a dependency on multiple controlled substances that should not
5 have been prescribed to him given the lack of medical justification or medical indication for
6 inappropriate prescribing of such medications by Respondent.

7 PATIENT 4

8 62. Patient 4 was an adult female patient with a history of anxiety and back pain.
9 Respondent prescribed the following medications to Patient 4 on a consistent basis over the
10 period of approximately 2016 to approximately 2018: clonazepam and hydrocodone.

11 63. The standard of care under such circumstances requires a medical history and
12 physical exam, which includes an assessment of the patient's pain, including physical and
13 psychological status and function; substance abuse history; history of prior pain treatments, and
14 assessment of any other underlying or co-existing conditions. Finally, it should include
15 documentation of recognized medical indications for the use of controlled substances. A history
16 and physical was not documented in any clinic visits by Respondent. Medical records for Patient
17 4 for the period between approximately January 1, 2016 through approximately June 30, 2018,
18 provide no evidence that a substance abuse history was performed. The records are consistently
19 illegible and lack documentation of clearly recognized medical indications for the use of
20 controlled substances given the patient's medical conditions. The records also show a lack of a
21 review of systems and the lack of a physical examination. There was no clear indication for
22 Respondent's prescribing of controlled substances for treatment of Patient 4's known medical
23 conditions.

24 64. The lack of a documented substance abuse history and lack of a review of systems
25 and physical examination for Patient 4 constitute an extreme departure from the standard of care.
26 The prescribing of controlled substances without a clear medical indication for the controlled
27 substance also is an extreme departure from the standard of care.

1 65. The standard of care requires that the medical records contain stated objectives that
2 may include relief of pain or relief of the medical condition requiring controlled substances
3 and/or improved physical or psychological function or ability to perform certain tasks or activities
4 of daily living. This should also include any plans for further diagnostic evaluations and
5 treatments, such as a rehabilitation program. From approximately January 1, 2016 through
6 approximately June 30, 2018, the medical records from the period reflect repeated departures
7 from the standard of care given Respondent's failure to document a specific treatment plan for
8 this patient. The medical records did not have a clear plan or objectives for the ongoing
9 prescribing of controlled substances.

10 66. Respondent's failure to document a specific treatment plan for this patient constitutes
11 repeated extreme departures from the standard of care. There was no evidence to support that
12 Respondent ordered any additional diagnostic evaluations or treatment for the patient related to
13 the indications for prescribing the numerous controlled substances.

14 67. Under such circumstances, the standard of care requires that the medical records
15 document that the physician discussed the risks and benefits of using controlled substances and
16 other treatment modalities. An actual written consent is not required but is recommended. There
17 was no evidence in the medical records that Respondent discussed potential side effects and risks
18 of controlled substances. There were no written consent forms noted in the medical records.
19 Respondent's clinical notes do not indicate that the risks of controlled substances were discussed.
20 Failure to discuss the potential side effects and risks of ongoing prescribing of controlled
21 substances with the patient constitutes an extreme departure from the standard of care.

22 68. The standard of care requires the medical records reflect that the physician is
23 periodically reviewing the course of treatment and prescribing controlled substances for the
24 patient, and making appropriate modifications in treatment based on the patient's progress or lack
25 of progress. The medical records for Patient 4 fail to demonstrate that Respondent ever
26 performed a periodic review on the patient's ongoing treatment with controlled substances even
27 though the patient was prescribed multiple controlled substances over a period of over
28 approximately two years. There were multiple extreme departures from the standard of care over

1 a period of approximately two years for failure to perform periodic reviews of the patient's
2 treatment and status in the setting of repeated prescribing and refilling of multiple controlled
3 substances.

4 69. The standard of care requires the physician consider obtaining additional evaluations
5 and consultations, especially with complex pain problems. Special attention should be given to
6 patients who are at risk for misusing their medication or have a history of drug addiction or
7 substance abuse. Such patients require extra care and monitoring and documentation, and
8 consultation with an addiction medicine specialist and, if indicated, a pain management specialist.
9 There is no evidence in the medical records that Respondent had made any referrals or obtained a
10 consultation related to prescribing the controlled substances for this patient. There is no evidence
11 that Respondent assessed the patient for any prior history of drug addiction or substance abuse.

12 70. Respondent's failure to obtain a consultation in this patient was a simple departure
13 from the standard of care.

14 71. The standard of care requires the physician to maintain accurate and complete
15 records, demonstrating a history and exam along with evaluations and consultations, treatment
16 plans and objectives, informed consent, medications prescribed, and periodic review
17 documentation. The medical records for all visits by Patient 4 are largely illegible in all sections
18 of the progress note. Likewise, the review of systems and physical examination sections of the
19 progress notes were largely illegible, making it impossible to determine why the patient was
20 being seen, what part of the physical examination was performed and what the specific
21 assessment and plan of care was for the patient. The records lacked medical indication for the
22 medications prescribed over a period of over approximately two years.

23 72. Respondent's failure to document standard guidelines in the use of controlled
24 substances for Patient 4 was an extreme departure from the standard of care.

25 73. Respondent's treatment of Patient 4 caused harm. The inappropriate prescribing of
26 controlled substances (in this case, clonazepam and hydrocodone) by Respondent without proper
27 justification or medical indication for such substances over a period of approximately three years
28 led to placing Patient 4 at an unnecessarily increased risk for significant morbidity and mortality

1 and potential harm given her pre-existing chronic medical conditions. Patient 4 had a history of
2 anxiety and back pain. She was prescribed multiple controlled substances by Respondent that had
3 high potential for abuse and dependency in a patient with multiple medical conditions who is at
4 risk for exacerbation of said co-morbidities by taking unnecessarily prescribed controlled
5 medications. This resulted in Patient 4 developing a dependency on multiple controlled
6 substances that were prescribed without medical indication. Patient 4 unnecessarily developed a
7 dependency on multiple controlled substances, which should not have been prescribed to her
8 given the lack of medical justification or medical indication for inappropriately prescribing of
9 such medications by Respondent.

10 **FIRST CAUSE FOR DISCIPLINE**

11 (Gross Negligence)

12 74. By reason of the facts set forth in paragraphs 16 through 30 (Patient 1), 31 through 47
13 (Patient 2), 48 through 61 (Patient 3), and 62 through 73 (Patient 4) above, Respondent is subject
14 to disciplinary action under Code section 2234, subdivision (b), in that he committed gross
15 negligence in his care and treatment of Patients 1, 2, 3, and 4 as follows:

16 75. The facts and allegations in paragraphs 16 through 73, above, are incorporated by
17 reference and re-alleged as if fully set forth herein.

18 **SECOND CAUSE FOR DISCIPLINE**

19 (Repeated Negligent Acts)

20 76. By reason of the facts set forth in paragraphs 16 through 30 (Patient 1), 31 through 47
21 (Patient 2), 48 through 61 (Patient 3), and 62 through 73 (Patient 4) above, Respondent is subject
22 to disciplinary action under Code section 2234, subdivision (c), in that he committed repeated
23 negligent acts in his care and treatment of Patients 1, 2, 3, and 4, as follows:

24 77. The facts and allegations in paragraphs 16 through 73, above, are incorporated by
25 reference and re-alleged as if fully set forth herein.

26 **THIRD CAUSE FOR DISCIPLINE**

27 (Failure to Maintain Adequate and Accurate Records)
28

1 78. By reason of the facts set forth in paragraphs 16 through 30 (Patient 1), 31 through 47
2 (Patient 2), 48 through 61 (Patient 3), and 62 through 73 (Patient 4) above, Respondent is subject
3 to disciplinary action under Code section 2266 for failure to maintain adequate and accurate
4 records of patient care in his care and treatment of Patients 1, 2, 3, and 4.

5 79. The facts and allegations in paragraphs 16 through 73, above, are incorporated by
6 reference and re-alleged as if fully set forth herein.

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8 **FOURTH CAUSE FOR DISCIPLINE**

9 (Prescribing Without an Appropriate Prior Examination and Medical Indication)

10 80. By reason of the facts set forth in paragraphs 16 through 30 (Patient 1), 31 through 47
11 (Patient 2), 48 through 61 (Patient 3), and 62 through 73 (Patient 4) above, Respondent is subject
12 to disciplinary action under Code section 2242 for prescribing controlled substances without an
13 appropriate prior examination and medical indication to Patients 1, 2, 3, and 4.

14 81. The facts and allegations in paragraphs 16 through 73, above, are incorporated by
15 reference and re-alleged as if fully set forth herein.

16 **FIFTH CAUSE FOR DISCIPLINE**

17 (Repeated Acts of Clearly Excessive Prescribing of or Administering of Drugs)

18 82. By reason of the facts set forth in paragraphs 16 through 30 (Patient 1), 31 through 47
19 (Patient 2), 48 through 61 (Patient 3), and 62 through 73 (Patient 4) above, Respondent is subject
20 to disciplinary action under Code section 725 for repeated acts of clearly excessive prescribing or
21 administration of drugs to Patients 1, 2, 3, and 4.

22 83. The facts and allegations in paragraphs 16 through 73, above, are incorporated by
23 reference and re-alleged as if fully set forth herein.

24 **SIXTH CAUSE FOR DISCIPLINE**

25 (Unprofessional Conduct)

26 84. By reason of the facts set forth in paragraphs 16 through 83, above, Respondent is
27 subject to disciplinary action under Code section 2234 for unprofessional conduct in his care and
28 treatment of Patients 1, 2, 3, and 4.

85. The facts and allegations in paragraphs 16 through 83, above, are incorporated by reference and re-alleged as if fully set forth herein.

PRAYER

WHEREFORE, Complainant requests that a hearing be held on the matters herein alleged, and that following the hearing, the Medical Board of California issue a decision:

1. Revoking or suspending Physician's and Surgeon's Certificate Number A 46373, issued to Respondent Moshen T. Moghaddam, M.D.;
2. Revoking, suspending or denying approval of his authority to supervise physician assistants and advanced practice nurses;
3. If placed on probation, ordering him to disclose the disciplinary order to patients pursuant to section 2228.1 of the Code;
4. If placed on probation, ordering him to pay the Board the costs of probation monitoring; and
5. Taking such other and further action as deemed necessary and proper.

DATED: MAR 24 2021

WILLIAM PRASIFKA
Executive Director
Medical Board of California
Department of Consumer Affairs
State of California

Complainant:

LA2021600806